ABSTRACT

A 1,3-dihydro-2H-indol-2-one derivative expressed by Formula 1 (wherein R_1 is a halogen atom, a C_1 to C_4 alkyl group, etc., and R_2 is a hydrogen atom, a halogen atom, etc., or R_2 is in the 6-position of the indol-2-one and R_1 and R_2 join together to form a C_3 to C_6 alkylene group, R_3 is a halogen atom, a hydroxyl group, etc., and R_4 is a hydrogen atom, a halogen atom, a C_1 to C_4 alkyl group, etc., or R_4 is in the 3-position of the phenyl and R_3 and R_4 join together to form a methylenedioxy group, R_5 is a hydrogen atom or a fluorine atom, R_6 is an ethylamino group, a dimethylamino group, etc., R_7 is a C_1 to C_4 alkoxy group, and R_8 is a C_1 to C_4 alkoxy group), or a pharmaceutically acceptable salt of this derivative. This is a novel compound that has antagonistic activity against an aruginine-vasopressin V1b receptor.